



A59

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>4</sup> :  A61K 39/39, C07J 17/00, 63/00		A1	(11) International Publication Number: WO 90/03184  (43) International Publication Date: 5 April 1990 (05.04.90)									
<p>(21) International Application Number: PCT/SE89/00528</p> <p>(22) International Filing Date: 28 September 1989 (28.09.89)</p> <p>(30) Priority data:</p> <table> <tr><td>251,576</td><td>30 September 1988 (30.09.88)</td><td>US</td></tr> <tr><td>8901027-6</td><td>22 March 1989 (22.03.89)</td><td>SE</td></tr> <tr><td>8902780-9</td><td>16 August 1989 (16.08.89)</td><td>SE</td></tr> </table> <p>(71)(72) Applicants and Inventors: MOREIN, Bror [SE/SE]; Olinstigen 3, Vreta, S-755 90 Uppsala (SE), LÖVGREN, Karin [SE/SE]; Lindsbergsgatan 8C, S-752 40 Uppsala (SE). DALSGAARD, Kristian [DK/DK]; Ny Vordingborgvej 80, DK-4771 Kalvehave (DK). THURIN, Jan [SE/US]; 28 University News, Philadelphia, PA 19104-4756 (US). SUNDQUIST, Bo [SE/SE]; Bellmansgatan 30, S-754 28 Uppsala (SE).</p> <p>(74) Agents: HJÄRNE, Per-Urban et al.; H. Albihns Patentbyrå AB, Box 3137, S-103 62 Stockholm (SE).</p>		251,576	30 September 1988 (30.09.88)	US	8901027-6	22 March 1989 (22.03.89)	SE	8902780-9	16 August 1989 (16.08.89)	SE	<p>(81) Designated States: AT (European patent), AU, BE (European patent), CH (European patent), DE (European patent), DK, FI, FR (European patent), GB (European patent), HU, IT (European patent), JP, LU (European patent), NL (European patent), NO, SE (European patent), SU, US.</p> <p>Published <i>With international search report.</i></p>	
251,576	30 September 1988 (30.09.88)	US										
8901027-6	22 March 1989 (22.03.89)	SE										
8902780-9	16 August 1989 (16.08.89)	SE										
<p>(54) Title: MATRIX WITH IMMUNOMODULATING ACTIVITY</p> <p>(57) Abstract</p> <p>The invention claims an iscom matrix which is not a lipid vesicle comprising at least one lipid and at least one saponin but no intentional antigenic determinants and optionally also adjuvants for use as an immunomodulating agent, medicines, vaccines, kits containing the matrix and new saponins, and a process for preparing the new saponins. The invention also concerns a process for preparing the matrix characterized in that at least one sterol is solubilized in a solvent or detergent, the saponin or saponins are added, the other adjuvants and lipids are optionally also added, whereafter the organic solvent or the detergent may be removed for example with dialysis, ultra filtration, gel filtration or electrophoresis. The sterol and saponin might also be solubilized in the lipids and/or adjuvants.</p>												

***FOR THE PURPOSES OF INFORMATION ONLY***

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	ES	Spain	MG	Madagascar
AU	Australia	FI	Finland	ML	Mali
BB	Barbados	FR	France	MR	Mauritania
BE	Belgium	GA	Gabon	MW	Malawi
BF	Burkina Faso	GB	United Kingdom	NL	Netherlands
BG	Bulgaria	HU	Hungary	NO	Norway
BJ	Benin	IT	Italy	RO	Romania
BR	Brazil	JP	Japan	SD	Sudan
CA	Canada	KP	Democratic People's Republic of Korea	SE	Sweden
CF	Central African Republic	KR	Republic of Korea	SN	Senegal
CG	Congo	LI	Liechtenstein	SU	Soviet Union
CH	Switzerland	LK	Sri Lanka	TD	Chad
CM	Cameroon	LU	Luxembourg	TG	Togo
DE	Germany, Federal Republic of	MC	Monaco	US	United States of America
DK	Denmark				

Matrix with immunomodulating activity

The present invention concerns an iscom matrix comprising at least one lipid and at least one saponin with immunomodulating effect, a process for preparing the matrix, a vaccine and a kit comprising the same and new saponins for incorporation in the matrix and a process for preparing the new saponins.

Many microbial and viral antigens can be produced by modern techniques today. Their full promise in vaccines will however not be realized unless they are administered along with an effective adjuvant, an agent that increases antibody and/or cell-mediated immune responses.

The only adjuvants currently authorized for human use in most countries are aluminium hydroxide and aluminum phosphate which have been used for many years to increase antibody responses to e.g. diphtheria and tetanus toxoids. Although these adjuvants are sufficient for many vaccines, studies have shown that other adjuvants, e.g. Freund's complete adjuvant (FCA), and Quil A often are more efficacious in eliciting antibody response and cell-mediated immunity in experimental animals. In fact, they are frequently required for protection. However, FCA produces granulomas at injection sites, which makes them unacceptable for human and veterinary vaccines. In fact, even aluminiumhydroxide may give rise to reactions in form of granuloma at the injection site. For these reasons, many attempts are made to develop adjuvants with the efficacy of FCA but without undesirable side effects.

In Morein's EPC Patent Applications Nos. 83850273.0 and 30 85850326.1, there are described immunogenic complexes between antigenic determinants with hydrophobic regions and glycosides, among them triterpensaponins and especially Quil A, so called iscom complexes. In such an iscom, the amount of Quil A

can be about 10-100 times lower and produce the same antigenic effect as when Quil A in free form is mixed with the antigen.

European Patent Application 87200035.1 indicates that the presence of antigen is not necessary for formation of the basic iscom structure, this being possible to form from a sterol, such as cholesterol, a phospholipid, such as phosphatidylethanolamine, and a glycoside such as Quil A.

It has now been discovered that a phospholipid is not needed for the preparation of the basic iscom structure including no antigen. Instead a sterol, such as cholesterol in conjunction with a glycoside such as Quil A are the essential structural components assembled into a complex resembling the typical cage-like iscom structure, so called matrix. It has also turned out that the matrix has immunomodulating effects such as adjuvant or immunosuppressive effect.

The present invention concerns a complex between at least one lipid such as a sterol, preferably cholesterol, and one or more saponins, such as triterpensaponins, especially Quil A or subcomponents thereof which is not a lipid vesicle without any intentional antigens or antigenic determinants for use as an immunomodulating agent. Thus, there is not integrated any antigenic component as is done in an iscom. This matrix has adjuvant effect and can be used mixed together with one or more antigens preferably in multimeric form.

In this iscom matrix there is also possible to integrate other adjuvants with hydrophobic regions. Addition of other lipids may be required to facilitate the inclusion of other adjuvants. Thus the present invention also concerns a complex containing lipids and adjuvants, other than cholesterol and saponins. Such complexes contains the matrix consisting of cholesterol and saponin, preferably Quil A or subcomponents thereof, one or more other adjuvants and one or more lipids

other than cholesterol. These are preferably not lipid vesicles or liposomes and have a very special structure in electron microscopy.

5    Liposomes have been described in the literature and their general structure is well known to biological research workers. Liposomes are vesicles comprising one or more series of lipid layers forming onion-like structures spaced one from another by aqueous material.

10    The matrix can be injected in an animal or human being as a mixture with the antigen in multimeric form. Alternatively the matrix and the antigen can be injected separately. In this case the best results are obtained if the adjuvant matrix and  
15    the antigen are injected in regions which are drained into the same lymphatic gland. When the adjuvant is presented in multimeric form in a matrix according to the invention the dose of adjuvant may be lowered as compared with when the adjuvant is injected separately in monomeric form or in an  
20    undefined form. This implies that toxic side effects caused by adjuvants when used conventionally, e.i. when they are injected alone as such, can be lowered or avoided. The dose of adjuvant can, however, not be lowered as much as is done in  
25    the iscom complexes according to the above mentioned patent applications.

When an adjuvant is used in a matrix according to the invention, the antigen is not integrated in the same particle as the adjuvant as is done in an iscom particle according to the  
30    above mentioned EPC Patent Applications. This implies that one can use antigens without amphiphatic properties or antigens which can not be forced to expose hydrophobic regions. As an example it can be mentioned that some viruses do not have amphiphatic proteins, e.g. picornavirus, adenovirus or parvovirus,  
35    but they have a form of submicroscopic particle with the antigen presented in several copies, i.e. as multimers.

For such viruses it is more practical to inject them together with the new adjuvant complex than to couple hydrophobic groups to them or create hydrophobic groups by other means (e.g. partial denaturation) and integrate them into an iscom particle.

5

Typically, the present matrix contains sterol, preferably cholesterol, and one or more saponins in a molar ratio of about 1 to 1 or in a weight ratio of about 1 to 5. The complexes have an open spherical structure consisting of circular subunits or parts of the spherical structure revealed by electron microscopy. They have a sedimentation coefficient of about 20 S.

10

15 When other adjuvants are integrated, the lipid-adjuvant-matrix typically contains sterol and saponin in a molar ratio of about 1:1 and the other adjuvants and lipids together make up to about 1 molar. For such a matrix the molar ratio of sterol; saponin; other adjuvant and lipids is about 1:1:1. Thus the 20 molar ratio of sterol; saponin; other adjuvant and other lipids is 1:1:0,1-1; 0,1-1, i.e. additional lipid or adjuvant may be present in the matrix until its molar ratio (or the sum of their molar ratios) is a half that of the saponin and sterol present.

25

The structure as revealed by electron microscopy is the same as for iscom and matrix (see Figure 1).

30

The sedimentation coefficient, being dependent on the density of material incorporated into the matrix, is about 12-22 S for matrixes containing cholesterol, saponin, other adjuvants and lipids.

35

The saponins can be any saponin with hydrophobic regions such as those described in R Tschesche and Wulf, Chemie und Biologie der Saponine in Fortschritte der Chemie Organischer

Naturstoffe, published by W Herz, H. Grisebach, G W Kirby, Vol 30 (1973), especially the strongly polar saponins, primarily the polar triterpensaponins such as the polar acidic bisdesmosides, e.g. saponin extract from Quillajabark Aralo-

5 side A, Chikosetsusaponin IV, Calendula-Glycoside C, Chiko-ssetsusaponin V, Achyranthes-Saponin B, Calendula-Glycoside A, Araloside B, Araloside C, Putranjia-Saponin III, Bersama-saponiside, Putrajia-Saponin IV, Trichoside A, Trichoside B, Saponaside A, Trichoside C, Gypsoside, Nutanoside, Dianthoside

10 C, Saponaside D, preferably aescine from Aesculus hippocastanum (T Patt and W Winkler: Das therapeutisch wirksame Prinzip der Rosskastanie (Aesculus hippocastanum), Arzneimittelforschung 10(4), 273-275 (1960) or sapoalbin from Gyposophilla struthium (R Vochten, P Joos and R Ruyssen: Physico-chemical properties of sapoalbin and their relation to the foam stability, J Pharm Belg 42, 213-226 (1968), especially saponin extract from Quillaja saponaria Molina, primarily the DQ-extract which is produced according to K Dalsgaard: Saponin Adjuvants, Bull Off Int Epiz 77 (7-8), 1289-1295 (1972) and

15 Quil A which is produced according to K Dalsgaard: Saponin Adjuvants III, Archiv für die Gesamte Virusforschung 44, 243-254 (1974). Quil A and subfragments thereof are preferred, especially the fragments B2, B3 and B4B described below.

20

25 The present invention also provides new glycosylated triterpenoid saponins derived from Quillaja Saponaria Molina of Beta Amyrin type with 8-11 carbohydrate moieties which have the following characteristics:

30 a) Substance B2 has a molecular weight of 1988, a carbon 13 nuclear magnetic resonance (NMR) spectrum as indicated in Figures 5A and 6A and a proton NMR spectrum as shown in Figures 11A and 12A.

35 b) Substance B3 has a molecular weight of 2150 and has a carbon 13 NMR spectrum as shown in Figures 5B and 6B, and a proton NMR spectrum as shown in Figures 11B and 12B.

c) Substance B4B has a molecular weight of 1862, a carbon 13 NMR spectrum as shown in Figures 5C and 6C, and a proton NMR structure as shown in Figures 11C and 12C.

5 Compounds B2 and B3 have adjuvant activity in their own right. The present invention also relates therefore to the use of these compounds as adjuvants. Compound B4B is of use in the preparation of an iscom matrix. B2 and B3 having adjuvant activity can be included in the matrix.

10 Matrix can be produced from a sterol such as cholesterol and the saponin B4B. Such a matrix does not seem to have any potent adjuvant activity. In order to potentiate the adjuvant activity in this matrix, it is possible and even preferable to integrate the saponins B2 and/or B3 and/or any other substance with adjuvant effect and with hydrophobic groups. If the adjuvants do not contain any hydrophobic groups such groups might be coupled to them by use of known chemical methods. If other adjuvants than B2 or B3 are to be integrated, there are 15 preferably incorporated further lipids as listed on page 13, last paragraph and paragraphs 1-3 on page 14.

20 In the sterol-B4B matrix, it is also possible to integrate immunosuppressive substances containing hydrophobic groups or 25 to which such groups have been coupled.

It is also possible to use the sterol-B4B matrix as an immuno-modulating agent in mixture with adjuvants, immunosuppressive substances or antigens or mixtures thereof.

30 As immunodulating agents are considered substances that enhance, suppress or change the immune system such as adjuvants, suppressors, interleukins, interferons or other cytokines.

35 The invention preferably concerns a matrix containing a sterol, especially cholesterol, B4B and either of B2 and B3 or

both. When matrix is prepared from cholesterol and Quil A, it comprises B2, B3 and B4B.

The matrixes can be produced by solubilizing at least one  
5 sterol in a solvent, adding the saponin or saponins, and  
possibly the other adjuvants and lipids, whereafter the  
solvent might be removed and the matrix transformed into a  
solution where its components are not soluble, e.g. a water  
solution. This can be done with gel filtration, ultra filtra-  
10 tion, dialysis or electrophores. The matrices may then be  
purified from excess of sterol and Quil A e.g. by centrifuga-  
tion through a density gradient, or gel filtration. As solvent  
there might be used water or the solubilizing agents or deter-  
gents mentioned below.

15 The only limiting factor for matrix formation to take place is  
the time needed in different physico-chemical environments,  
the major rate limiting factor being the poor solubility of  
the sterol, e.g. cholesterol, in water, in which the matrix  
20 forming saponins are freely soluble.

Thus it has been shown that with Quil A and cholesterol even  
in solid phase matrix-like formation takes place after a  
relatively long time, e.g. about 1 month. Cholesterol must be  
25 brought into contact with Quil A or its purified components.  
If the cholesterol is brought into colloidal water suspension  
through treatment by ultrasonication and treatment by ultra-  
-turrax, matrix is formed with Quil A after about 12 hours.

30 Consequently, any other substance such as a detergent added to  
the water, and which will increase the solubility of chole-  
sterol in the aqueous medium, will decrease the time needed  
for the formation of matrix. It is thus possible to produce a  
matrix from cholesterol, water and Quil A or the subcomponents  
35 thereof, if the cholesterol is brought to a colloidal form. It  
is, however, more practical to add a detergent or a solvent.

Preferably the saponins are used from a concentration of at least their critical micelle formation concentration (CMC). For Quil A this implies a concentration of at least 0,03 % by weight.

5

As solubilizing agent there can be used detergents such as non-ionic, ionic i.e. cationic or anionic or zwitter-ionic detergent such as Zwittergent or detergent based on gallic acid which is used in excess. Typical examples of suitable 10 non-ionic detergents are N-alkanoyl-N-alkyl-glucamines, polyglycol esters and polyglycol ethers with aliphatic or aralyphatic acids and alcohols. Examples of these are alkylpolyoxyethylene ethers with the general formula  $C_n H_{2n+1} (OCH_2 CH_2)_x OH$ , shortened to  $C_n E_x$ ; alkyl-phenyl polyoxiethylene ethers containing a phenyl ring between the alkyl group and the polyoxyethylene chain, abbreviated 15  $C_n \phi E_x$ , e.g. Triton X-100 = tert.- $C_8 E_{9,6}$  (octyl-phenolether of polyethylene oxide), acylpolyoxyethylene esters; acylpolyoxyethylene sorbitane esters, abbreviated  $C_n$  20 sorbitane  $E_x$ , e.g. Tween 20, Tween 80,  $\beta$ -D-alkylglucosides, e.g.  $\beta$ -D-octylglucoside. Typical examples of suitable ionic detergents are gallic acid detergents such as e.g. cholic acid, desoxycholate, cholate and CTAB (cetyltrimmonium bromide). Even conjugated detergents such as e.g. taurodeoxy- 25 cholate, glycodeoxicholate and glycocholate can be used. Other possible solubilizing agents are lysolecithin and synthetic lysophosphoillipids. Even mixtures of the above-mentioned detergents can be used. When using the dialysis method the detergents should be dialysable in not too long time.

20

Some surface active substances greatly facilitate matrix formation. These include the intrinsic biological membrane lipids with a polar head group and a non-polar aliphatic chain e.g. phosphatidyl choline (negatively charged) and phosphatidyl ethanolamine (positively charged).

35

Solubilizing can also be performed with alcohols, organic solvents or small amphiphatic molecules such as heptane-1,2,3-triol, hexane-1,2,3-triol or caotrophic substances, acetic acid, such as trifluoro-acetic acid, trichloro-acetic acid, 5 urea or quanidine hydrochloride.

Preferably to be used are ethyl alcohol, dioxane, ether, chloroform, acetone, benzene, acetic acid, carbon disulphid, MEGA-10 (N-decanoyl-N-methyl glucamine) and  $\beta$ -octylglucoside.

10 Various yields of matrix can be obtained with these substances, and the overall picture is that more matrix is formed the higher the concentration of the detergent is in the system.

15 It is technically possible to produce, purify, and sterilize matrix in any of the systems described. Therefore the adjuvant active technical preparations of matrix may contain solubilizing agents if their chemical nature and their concentration is acceptable in the final product, e.g. for vaccine purposes.

20 However, in many cases it will be necessary to remove the solubilizing agent from the matrix by dialysis, ultrafiltration or column chromatographic techniques. It is even possible to dilute the preparation until an allowed concentration of a given solubilizing agent or detergent is reached. The preparation is diluted with water or a physiologically acceptable solution preferably to a concentration below the CMC for the solubilizing agent or detergent in the system (the preparation) used.

25 30 The solubilizing agent might be incorporated in the matrix in a molar ratio of sterol; saponin; further lipid adjuvants or solubilizing agent 1:1:1, i.e. molar ratio of the sum of lipid, adjuvants and solubilizing agent is up to half the molar that of saponin and sterol.

35 The solubilizing agent might alternatively be left mixed with the iscom matrix.

In order to be integrated the solubilizing agent and other immunomodulating components, should have at least one hydrophobic region. If not present such hydrophobic regions can be coupled to the components before the matrix is made.

5

Examples of adjuvants that can be incorporated in iscom matrix are any adjuvant, natural or synthetic, with desired immunomodulatory effect, e.g. muramyl dipeptide (MDP)-derivatives, such as fatty acid, substituted MDP, threonyl analogs of MDP; amphipatic copolymers, aliphatic amines such as avridine or DDA, poly anions such as Dextran sulphate, lipopolysaccharides such as saponins (other than Quil A). ("Future prospects for vaccine adjuvants", Warren, H.S. (1988) CRC Crit. Rev. Immunol. 8:2, 83-101; "Characterization of a nontoxic monophosphoryl lipid A", (1987) Johnson, A.G. et al, Rev. Infect. Dis. 9:5, 5512-5516; "Developmental status of synthetic immunomodulators", Berendt, M.J. et al (1985), Year Immunol. 193-201; "Immunopotentiating conjugates", Stewart-Tull, D.E., Vaccine, 85, 3:1, 40-44).

10

These four references are hereby incorporated as references.

15  
20  
25

The following zwitterionic, neutral, positive and negative detergents are examples of detergents that have immunomodulating, especially adjuvant activity:

30  
35

Nonionic block polymer surfactants containing hydrophilic polyoxyethylene (POE) and hydrophobic polyoxypropylene (POP) that differed in mol weight percentage of POE and mode of linkage POP to POE (BASF Wyandotte Corp.), such as L72, L81, L92, pluronic L101, L121, 2531 and 31R1; octablocks T1501; B-D-octylglucosid; cationic surfactants such as dimethyldioctadecylammonium bromide (DDA), octadecylamine (OCT), and cetyltrimethylammonium bromide (CTAB); maltostose tetrapalmitate, trehalose monomycolate, trehalose dibehenyl-behenate; zwittergent detergents (N-alkyl-N,N-dimethyl-

-ammonio-3-propanesulphonate) Z3-8, Z3-10, Z3-12, Z3-14, Z3-16, obtained from Calbiochem (La Jolla, CA, USA); Z3-18 obtained from Serva (Heidelberg, FRG), Myrij 45, Brij 52, Brij 58 (also from Serva), and dioctylsulphosuccinate and Tween 20, Tween 80, Triton X-100 and sodium deoxycholate.

These detergents can be used as both detergents and adjuvants and be incorporated in the iscom matrix.

The following are examples of immunosuppressive agents that can be incorporated in a sterol (preferably cholesterol) B4B matrix: cyclosporin A, diodecyl dimethyl ammonium bromide, cationic single chain amphiphiles with more than 10 carbon atoms and preferably more than 15 carbon atoms, double chain amphiphiles with up to 14 carbon atoms, preferably up to 12 carbon atoms.

In the case a desired adjuvant or immunosuppressive agent do not have suitable hydrophobic properties, it has to be modified to comprise a hydrophobic domain for incorporation into the matrix.

The hydrophobic group that can be coupled to non-hydrophobic adjuvants are straight, branched, saturated or unsaturated aliphatic chains having 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24 and 25 carbon atoms, such as lipids, preferably 6, 7 and 8 carbon atoms; small peptides with 1, 2, 3, 4 or 5 amino acids, preferably 2, 3, 4, selected from Trp, Ile, Phe, Pro, Tyr, Leu, Val, especially Tyr; choline acid, ursodesoxycholine acid or cholesterol derivatives.

These hydrophobic groups must be bonded to a group that can be coupled to the non-hydrophobic protein such as carboxyl-, amino-, disulphide-, hydroxyl-, sulphhydryl- and carbonyl group, such as aldehyde groups.

As hydrophobic groups that can be coupled are selected preferably carboxyl, aldehyde, amino, hydroxyl, and disulphide derivatives of methane, ethane, propane, butane, hexane, heptane, octane and peptides containing Cys, Asp, Glu, Lys,

5 preferably octanal and Tyr.Tyr-Cys, -Asp or -Glu. The hydrophobic groups with a group that can be coupled must be dissolved in water with the aid of for example the solubilizing agents and detergents mentioned above or hydrochloric acid, acetic acid, 67% by volume acetic acid, caustic liquor, 10 ammonia, depending on what substance is to be dissolved. pH is then adjusted to the neutral direction without the substance precipitating; here it is to make sure that there is not obtained a pH-value that denaturates the protein to which the hydrophobic group is to be coupled.

15

Hydrophobic groups with a carboxyl group as coupling molecule can be coupled to the adjuvants through water-soluble carbodiimides or composite anhydrides. In the first case the carboxyl group is activated at pH 5 with carbodiimide and mixed with 20 the protein dissolved in buffer pH 8 with a high phosphate content. In the latter case the carboxy compound is reacted with isobutylchloroformate in the presence of triethylamine in dioxane or acetonitrile, and the resulting anhydride is added to the protein at pH 8 to 9. It is also possible to convert the 25 carboxyl group with hydrazine to hydrazide which together with aldehydes and ketones in periodate-oxidized sugar units in the protein gives hydrazone bonds.

20

The amino groups with nitrous acid can at low temperature be converted to diazonium salts, which gives azo bonds with Tyr, His and Lys.

35

The hydroxyl groups with succinic anhydride can be converted to hemisuccinate derivatives which can be coupled as carboxyl groups.

Aldehyde groups can be reacted with amino groups in the protein to a Schiff's base.

Several coupling groups and methods are described in Journal of Immunological Methods, 59 (1983) 129-143, 289-299, Methods in Enzymology Vol 93 pp 280-33, and in Analytical Biochemistry 116, 402-407 (1981) which are here incorporated as references.

The lipids other than sterol can be fats or fat resembling substances such as triglycerides or mixed triglycerides containing fatty acids with up to 50 carbon acids such as saturated fatty acids with 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29 and 30 carbon atoms e.g. butyric acid, caproic acid, caprylic acid, capric acid, lauric acid, myristic acid, palmitic acid, stearic acid, arachidic acid, behenic acid, lignoceric acid, or unsaturated fatty acids with up to 30 carbon atoms, such as hexadecene acid, oleic acid, linoleic acid, linolenic acid, arachidonic acid; hydroxy-fatty acids such as 9,10-dihydroxy stearic acid; unsaturated hydroxy fatty acids such as castor oil, branched fatty acids; glycerol ethers, waxes i.e. esters between higher fatty acids and monohydric alcohols; phospholipides such as derivatives of glycerol phosphates such as derivatives of phosphatidic acids i.e. lecithin, cephalin, inositol phosphatides, sphingosine derivatives with 14, 15, 16, 17, 18, 19 and 20 carbon atoms; glycolipids isoprenoids, sulpholipids, carotenoids, steroids, sterols, cholestanol, caprostanol, phytosterols, e.g. stigmasterol, sitosterol, myco sterols, e.g. ergosterol, bile acids e.g. cholic acid, deoxycholic acid, chenodeoxycholic acid, lithocholic acid, steroid glycosides, esters of vitamine A, or mixtures thereof.

These and other useful lipids are described in: Lipid biochemistry and introduction, Ed. M.I. Gurr, A.I. James, 1980, Chapman and Hall, London, New York, University Press Cambridge, which hereby is incorporated as a reference.

Preferably cholesterol phosphatidylcholine, liposomes or intralipid<sup>®</sup> (Oleum soya fractionate 200 g, Lechitinum fractionate vitello ovi 12 g, glycerol 22.5 g, and H<sub>2</sub>O up to 1 liter) are used.

5

The lipids can be added at any stage in the process, preferably before the addition of the saponin but lipids could also be added after the saponin.

10 The matrix is best produced by the dialysis method as follows.

Cholesterol dissolved in 20% MEGA-10 or any other suitable detergent, preferably a detergent that can be removed by dialysis, e.g. β-octylglucoside, (in H<sub>2</sub>O or a suitable buffer) is mixed with 5 times as much Quil A (solid or dissolved in water or a suitable buffer, e.g. PBS). The mixture is dialysed extensively against PBS, first over night at room temperature (because MEGA-10 will precipitate at +4°C), then at +4°C. The matrixes are purified from excess Quil A and cholesterol by pelleting through e.g. 30% (w/w) sucrose (e.g., a TST 41.13 rotor 18 h, 39.000 rpm, 10°C). The pelleted matrixes are dissolved in PBS (or any other suitable buffer) and the concentration adjusted to 1 mg/ml.

25 The present matrix can be used as an immunomodulating substance. It can be used as a potentiating agent for an immuno-suppressive substance or an adjuvant, either mixed therewith or integrated in the matrix.

30 A matrix containing a sterol such as cholesterol, saponins, adjuvants and optionally further lipids can be used as an adjuvant. It can be used for potentiating the antigenic effect of any antigen or antigenic determinants from any pathogenic organism or any fragments or subunits of, or derived from 35 these. Thus it can be used as an adjuvant for those antigens that are integrated in an iscom. Such antigens are mentioned

in the EPC-patent applications 83850273.0 and 85850326.1, which are hereby incorporated as references. Thus the matrix can be used as adjuvants together with antigenes or antigenic determinants derived from viruses with or without envelope, 5 bacteria, protozoa, mycoplasmas, helminths, mollusca or together with such whole organisms. The antigenes or antigenic determinants might further be hormones, enzymes, carbohydrates and carbohydrate-containing structures such as lipopolysaccharides, peptides or proteins or recombinants thereof.

10

The present invention thus also covers human or veterinary medicine, characterized in that it comprises at least one matrix and one or more antigenic or immunosuppressive substances and a pharmaceutically acceptable vehicle in mixture 15 or in separate compartments.

The invention also concerns a vaccine comprising an matrix, one or more antigens and a pharmaceutically acceptable vehicle.

20

Further the invention concerns a kit comprising such a medicine or vaccine.

25

In some medicines or vaccines the detergent used when preparing the matrix can be present if the detergent is allowed for the product in question.

The effect of the new adjuvant complex according to the invention will now be described in immunostimulating experiments.

30

1. Comparison between the immunogenic effects from antigens presented as iscoms, micelles or micellas plus the new matrix.

35

Mice were immunized with envelope protein from influenza virus in the form of iscom complex, micelles and micellas together with the new complex according to the invention (so called

matrix). The immune response was evaluated by measuring the antibodies with ELISA technique 15, 30, 44 and 50 days after injection. The following injections were made:

5    1. 5 µg Micell + 0,1 µg matrix were mixed and injected in the left foreleg.

2. 5 µg Micell + 0,1 µg matrix injected separately in the right and left foreleg respectively.

10

3. 5 µg Micell

4. 5 µg iscom prepared according to EPC 83850273.0

15

TABLE 1

<u>DAY</u>	<u>1</u>	<u>2</u>	<u>3</u>	<u>4</u>
15	23.700 ± 9.500	12.900 ± 14.500	18.900 ± 9.500	41.600 ± 1.000
20	30.800 ± 10.500	8.800 ± 7.100	9.800 ± 3.600	80.700 ± 21.700
44	30.000 ± 17.600	32.600 ± 17.300	17.900 ± 4.200	129.100 ± 78.400
50	309.300 ± 89.000	136.700 ± 103.700	87.600 ± 18.200	880.430 ± 295.500

No side effects in the form of local reactions were noted.

From this experiment one can conclude that envelope protein from influenza in the form of iscom or micelles plus matrix gives the highest antibody titres. Matrix can be presented in a very low dose and still have adjuvant effect. In order to get an adjuvant effect in mice, Quil A in free form is required in a dose a 100 times the dose of matrix, i.e. 30 10 µg. With that dose Quil A begins to give local side reactions. On order for matrix to have an obvious adjuvant

effect the antigen in multimeric form should be injected in the same region e.g. leg as the matrix, i.e. the injected adjuvant matrix complex and antigen should be presented in a region, that is drained to the same lymphatic gland.

5

2. Comparison between the immunogenic effects from envelope protein from influenza in the form of iscom or micelle with or without matrix or difteritoxoid (DT).

10 Mice were injected with envelope protein in the following forms:

1. 5 µg Iscom + 5 µg DT

15 2. 5 µg Iscom

3. 5 µg micelle

4. 5 µg micelle + 0,1 µg matrix

20

The antibody response in envelope proteins was estimated in the serum with ELISA-technique. The following results were obtained:

25

TABLE 2

<u>DAY</u>	<u>1</u>	<u>2</u>	<u>3</u>	<u>4</u>
15	52.800 1)	48.300	8.400	29.000
30	119.202	155.567	22.107	87.000
50	110.600	136.200	33.400	96.500
30	1.691.000	3.783.000	283.300	1.149.000
	562.800	2.529.000	512.300	976.500

1) ELISA-titer where the last dilution gives a significant positive value at 450 nm.

35

No visible side effects in form of local reactions could be noted.

One can conclude that envelope protein from influenza virus in the form of iscom or micelles plus the adjuvant complex (matrix) according to the invention gives the highest antibody titres. The dose of matrix can be kept very low, i.e.

5 0,1 µg, and still has a notable adjuvant effect.

3. Comparison between the immunogenic effects from diphtheria toxoid (DT) in monomeric form, monomeric DT + iscom containing envelope protein from influenza virus, monomeric DT in mixture 10 with Quil A and cholesterol and monomeric DT + adjuvant complex (matrix) according to the invention.

In this experiment diphtheria toxoid is used as a model antigen in monomeric form.

15

Mice were injected with diphtheria toxoid in the following forms:

1. 5 µg DT (diphtheria toxoid)

20

2. 5 µg DT + 5 µg iscom

3. 5 µg DT+ 0,5 µg Quil A + 0,1 µg CL (cholesterol)

25

4. 5 ug DT + 0,1 ug matrix

TABLE 3

DAY	1	2	3	4
30	-	-	-	-
15	≤ 30	≤ 30	≤ 30	≤ 30
30	≤ 30	≤ 30	≤ 30	≤ 30
50	≤ 30	≤ 30	≤ 30	≤ 30
65	≤ 30	90	≤ 30	10.000
80	≤ 30	60	90	1.100

35

The immungenic response to DT is low in all the groups. The

best result is obtained with mice immunized with diphtheria toxoid plus matrix according to the invention.

From the experiments above one can conclude that the best  
5 results are obtained when the matrix according to the invention is used together with the antigen in multimeric form. The matrix according to the invention has thus proved to give very good results as adjuvant compared with e.g. Quil A in free form. Thus it is worth noting that Quil A is effective as  
10 adjuvant in free form in doses such as 10 µg for mice, 50 µg for guinea-pigs and 1 mg for cattles. A practical volume for injection of a vaccine is 1 ml for small animals and 2 to 5 or 10 ml for big animals. As CMC (the critical  
15 micelle concentration) for Quil A is 0,03%, 1 ml will imply an amount of 300 µg when 1 ml is injected. After injection, however, due to the dilution effect, the concentration will become lower than CMC and the micelle will become unstable.

According to the present invention, however, the saponin and  
20 especially the Quil A molecules will be bounded together with cholesterol molecules so that a relatively stable complex is formed at very low concentrations. This complex is effective as adjuvant in a dose, which corresponds to 0,1 µg Quil A,  
i.e. 100 times lower than when Quil A is presented in free  
25 form.

The Figures show:

Fig. 1 shows an electron microscope picture of a typical matrix;

30 Fig. 2 shows U.V. eluation profiles for subfractions of Quil A;  
Fig. 3 demonstrates HPTLC-separation of Quil A and its sub-fractions;

Fig. 4 shows FAB-mass-spectra for the new substances according to the invention;

35 Fig.:s 5 and 6 show <sup>13</sup>C-NMR-spectra (2-regions) for the new substances;

Fig.:s 7, 8 and 9 show complete  $^{13}\text{C}$ -NMR-spectra for B2, B3 and B4B, respectively;

Fig. 10 shows the  $\beta$ -amyrin-skeleton;

Fig.:s 11 and 12 show the  $^1\text{H}$  NMR-spectra for the new substances;

Fig. 13 shows parts of the spectra in Fig.:s 7 and 8; and Fig. 14 shows a 2-dimensional NMR-spectrum for substance B3.

The invention will now be described further with the following example.

Example 1:

Matrix (Cholesterol-Quil A complex)

1 mg of cholesterol dissolved in 20 % MEGA-10 (in  $\text{H}_2\text{O}$ ) was mixed with 5 mg of solid Quil A. The Quil A was dissolved and the mixture was dialysed extensively against PBS, first over night at room temperature, then at +4°C. The iscom matrixes were purified from excess Quil A and cholesterol by pelleting through 30 % (w/w) sucrose (TST 41.13 rotor 18 h, 39.000 rpm, 10°C). The pelleted matrixes were dissolved in PBS and the concentration adjusted to 1 mg/ml (traced by a small amount of  $^3\text{H}$ -cholesterol).

Example 2:

MDP (muramyldipeptide, Sigma, adjuvant peptide) was conjugated to phosphatidyl ethanolamine (PEA) using N-ethyl-N'-(dimethylaminopropyl) carbodiimide hydrochloride as described by Lefrancier et al., 1977 (Lefrancier, P., Choay, J., Derrien, M. and Lederman, I. (1977) Int. J. peptide Protein Res. 9:249-257).

To 1 mg of cholesterol (in 20% MEGA-10 in  $\text{H}_2\text{O}$ ) was added an equimolar amount of MDP-PEA (in MEGA-10 or DMSO or any other water miscible solvent), an equimolar amount of phosphatidyl choline and 7 mg of Quil A (a slight excess in comparison to 5 mg that is required for IM-formation). After a short in-

cubation at room temperature (15-30 min) the mixture was extensively dialysed against PBS (room temperature 4-12 h, then at +4°C).

5 After completed dialysis, the matrix-complexes with the additional adjuvant integrated were purified from excess Quil A by pelleting through 10% sucrose.

Example 3:

10 To 1 mg of cholesterol (in 20% MEGA-10 in H<sub>2</sub>O) was added an equimolar amount of Avridine (N,N-dioctadecyl-N'N'-bis(2-hydroxyethyl)propenediamine (in MEGA-10 or DMSO or any other water miscible solvent), an equimolar amount of phosphatidyl choline and 7 mg of Quil A (a slight excess in comparison to 15 5 mg that is required for IM-formation). After a short incubation at room temperature (15-30 min) the mixture was extensively dialysed against PBS (room temperature 4-12 h, then at +4°C).

20 After completed dialysis, the matrix-complexes with the additional adjuvant integrated were purified from excess Quil A and adjuvant by pelleting through 10% sucrose (the same method as described on page 14, last paragraph).

25 Example 4:

To 1 mg of cholesterol (in 20% MEGA-10 in H<sub>2</sub>O) was added an equimolar amount of DDA (dimethyl dioctadecyl ammonium bromide (in MEGA-10 or DMSO or any other water miscible solvent), an equimolar amount of phosphatidyl choline and 7 mg of Quil A (a 30 slight excess in comparison to 5 mg that is required for matrix-formation). After a short incubation at room temperature (15-30 min) the mixture was extensively dialysed against PBS (room temperature 4-12 h, then at +4°C).

35 After completed dialysis, the matrix-complexes with the additional adjuvant integrated were purified from excess Quil A

and adjuvant by pelletting through 10% sucrose (the same method as described on page 14, last paragraph).

Example 5:

5 2 g of Mega 10 is added to 10 ml of water before the addition of 200 mg cholesterol, and the cholesterol is dispersed by ultrasonication/ultraturrax. As much as 1.6 ml of this mixture can be added to the 10 ml of 2% Quil A-solution. The reaction mixture clarifies completely after less than one hour indicating that all the cholesterol has been reacted. It can be seen in the electron microscope that the concentration of matrix is very high even if the concentration of detergent in this case is 10%. Removal of the detergent by dialysis or ultrafiltration does not quantitatively affect the number of matrix particles, and the solution of matrix stays completely clear.

10 This experiment indicates that matrix formation takes place when the surfactants are present in the reaction mixtures, and that complete matrix formation takes place in very high concentrations of detergent.

Example 6:

Preparation of the Quil A components B2, B3 and B4B according to the invention.

20 25 5 g Cortex quillajae (Nordiske Droege of Kemikalieforretning, Copenhagen, Batch nr 8372) and 50 ml destillated water was mixed by a magnetical stirrer for 3 hours at room temperature. The liquid phase was separated through a Büchner funnel by a filter paper and was purified by filtering through a Metricel 30 Gelman membrane 0.22 µ. Such an extract contains 2.5% dry material.

35 The crude extract was dialysed against 200 volumes of destillated water in a Visking-tube without weld 20/32 for 48 hours with exchange of water after 24 hours. This extract is called DQ.

The dialysed extract above was subjected to ion exchange chromatography. A column of DEAE-cellulose equilibrated with 0.1M Tris-HCl pH 7.5 was prepared (Whatman DE52) in a K 9/15 column (Pharmacia Fine Chemicals). The bed material was equilibrated with 0.1M Tris-HCl buffer pH 7.5. The column was eluted either stepwise or by a linear salt gradient at a flow rate of 60 ml/h using a peristaltic pump. 50 ml DQ was introduced on the column and 300 drop (equivalent to approx. 5 ml) fractions were collected. Under these conditions, some of the substances in DQ passed unbound through the column, as will be seen from Fig. 2A (peak A). Elution was continued until no UV absorption was detectable. The absorption of the effluent liquid was recorded at 280 nm by a Uvicord II system (LKB-Produkter), and fractions were collected by a Golden Retriever (ISCO). At this point a buffer containing 0.2M NaCl made up in start buffer was introduced. As can be seen in Fig. 2A, a peak B is eluted. However, some substances were still attached to the bed material to such a degree that elution was difficult even with concentrated NaCl. These substances were the ones that contributed to the brownish colour of DQ, whereas peak A and B were only slightly coloured or completely colourless, respectively. In the next purification step, peak B was pooled and subjected to gel exclusion chromatography on Sephadex G50 fine equilibrated with M/50 phosphate buffer pH 7.5 in a K 16/70 column eluted at a flow rate of 10 ml/h. Desalting was carried out on Sephadex G25 medium in a K 16/40 column. Elution was carried out using a hydrostatic head of 50 cm. As will be seen from Fig. 2B, the UV profile showed 3 peaks. Peak C was eluted in the void volume (as determined by Blue Dextran 2000, Pharmacia Fine Chemicals) and peak E was eluted in the total volume of the column (also determined by potassium chromate). Peak D was well separated from peak C and E, but as can be seen in Fig. 2B, the presence of a shoulder indicated that peak D consisted of at least two substances.

Consequently, peak D was pooled and subjected to a new separation on DEAE-cellulose. The starting conditions were the same as in the first ion exchange experiment, and as a result all the material was adsorbed to the column. Elution was now continued with a linear NaCl gradient increasing from 0 to 1 molar (made up in the start buffer) in the course of 300 ml.

The result of this experiment is shown in Fig. 2C. Two peaks F and G appeared in the UV profile. F was clearly separated and a single substance (section 3.3), but peak G could not be isolated since it was contaminated with F. In order to investigate the homogeneity of peak F, it was pooled, desalted on Sephadex G25, and rechromatographed in an identical experiment. As can be seen from Fig. 2D, only one symmetrical peak (H) appeared at the position of peak F.

15

Anyone of the fractions (also the DQ-extract) can be further purified as follows.

Fraction H was lyophilized and dissolved in chloroform/methanol/H<sub>2</sub>O (60:40:9, v/v/v). 200 mg was then applied to an HPLC column (4.5x50 cm) packed with silicic acid, Iatrobeads RS-8060 (Iatron Labs, Tokyo, Japan). A pump speed of 5 ml/min was used for pumping a total of 2 L solvent, collecting 200 fractions of 10 ml with a gradient of chloroform/methanol/H<sub>2</sub>O (60:40:9 to 50:40:10). The fractions were analyzed with thin layer chromatography in the following way. 2 µl of every second fraction was analyzed by developing thin layer liquid chromatography plates (TLC-plates) (HPTLC, Merek, Bodman Chemicals, Gibbstown, NJ) in chloroform/methanol/0.2% CaCl<sub>2</sub> (50:40:10 v/v/v) and the glycosides were determined by being greencoloured with anisaldehyde reagent (acetic acid/sulphuric acid/paraanisaldehyde (98:2:1)). The Quil A starting material was used as a reference of the R<sub>f</sub>-value. (See Fig. 3, which shows a HPTLC-separation of: lane 1, Quil A (fraction H); lane 2, B1; lane 3, B2; lane 4; B3; lane 5, B4A; and lane 6, B4B).

Fractions that comigrate with B2, B3 and B4B having identical R<sub>f</sub>-values were pooled and analyzed for purity with TLC. These crude fractions usually must be chromatographed two times in order to become pure enough. Fractions B1 and B4A are 5 inactive and therefore are not separated further.

The thus enriched components were further purified on an HPLC column (21.2x250 mm) packed with 5 µ spherical silica particles (Zorbax Si, DuPont, Wilmington, DE). 40 mg enriched fraction B2, B3 or B4B dissolved in 1 ml chloroform/methanol/H<sub>2</sub>O (60;40;9 v/v/v) was put on the column. A pump speed of 3 ml/min was used for pumping a total of 0.9 l solvent, collecting 300 fractions of 3 ml with a gradient of chloroform/methanol/H<sub>2</sub>O (60;40;9 to 50;40;10 v/v/v). Fractions were 10 analyzed on glass-backed HPLTC-plates as above. Purified 15 fractions were pooled and evaporated to dryness in a rotary evaporator <30°C, dessicciated and stored in <-20°C. Approximately 20-25 rounds of this purification step was used i.e. 20 using (20-25)x200 mg = 4-5 g Quil A starting material, including rechromatography to prepare 1 g of fraction B3. The yield of B2 and B4B was about 40% of the yield of B3.

The so prepared components B2, B3 and B4B were analyzed as follows.

25

a) Mass Spectrometry

Negative FAB-MS, Fig. 4, and positive FAB-MS (data not shown) 30 were carried out for determination of molecular weights of the purified Quil A ocmponents B2, B3, B4A, and B4B. The data shown in Fig. 4 are preliminary and will have to be re-acquired in a neutral pH matrix such as glycerol rather than 35 in triethanolamine which was used for the spectra shown in Fig. 4. This is necessary because extreme alkali-lability of the compounds, pH > 8.5 have been demonstrated. Peaks at m/z 595, 744, and 893 stem from the matrix triethanolamine and

should be disregarded. Our fraction B4A, which does not have any adjuvant or ISCOM particle forming capacity, seem to be identical with that described by Komori et al (for structure, see Fig. 10). The peaks corresponding to molecular weights of 5 the three thus far most interesting glycosides are at: m/z 1988, B2; m/z 2150, B3; and m/z 1862, B4B.

b)  $^{13}\text{C}$ -NMR

10 Fig.:s 5 and 6 show two regions, aliphatic carbon (8-45 ppm) and anomeric carbon (90-115 ppm), respectively, of the  $^{13}\text{C}$ -NMR spectra for the full size fractions: A, B2 (20 mg); B, B3 (80 mg); and C, B4B (40 mg). All spectra were obtained in the solvent-system, chloroform/methanol/water (30:60:8, v/v/v). The triterpenoid region is well resolved (8-45 ppm, Fig. 6) and has been partially assigned as seen in Table 4.

TABLE 4

20 Partial  $^{13}\text{C}$ -NMR signal assignment (ppm) for  $\beta$ -amyrin five-ring segment of fractions B2, B3, and B4B (see Fig. 5) obtained in chloroform/methanol/water (30:60:8, v/v/v).

25

<u>Carbon#</u>	<u>B2</u>	<u>B3</u>	<u>B4B</u>	<u>Reference</u>
C9 <sup>a</sup>	-b	-b	-b	45.5
C10	36.4	36.2	36.3	37.0
C12	122.5	122.2	122.5	123.1
C13	144.1	146.6	143.6	144.8
C14	41.5	41.8	41.9	41.6
C15	30.0	30.5	30.7	30.7
C18	43.0	42.8	41.9	42.7
C20	30.7	30.6	30.6	30.7
C25	16.0	16.1	16.0	15.9
C26	17.7	17.3	17.6	17.6
C29	32.9	32.9	32.9	32.2

35

<sup>a</sup> Numbered as in Fig. 5.

<sup>b</sup> Hidden under methanol signal of solvent (confirmed with a DEPT experiment).

These assignments have been performed from studying a large number of reference-spectra obtained in various solvents and by analyzing the signals that are solvent-independent by a statistical comparison (data not shown). Fig. 6 shows the 5 region between 80-148 ppm in the spectra of the three compounds, A, B2; B, B3; and C, B4B, featuring two double-bond carbon signals at 122 and 143 ppm corresponding to C-12 and C-13, respectively, in the  $\beta$ -amyrin skeleton (Fig. 10). The 10 anomeric-carbon region, between 90-115 ppm, shows the presence of approximately 9-10 signals corresponding to the same amount of sugar-residues in the compounds.

Conclusion: Structural differences can be identified between fractions: A, B2; B, B3; and C, B4B, in both spectral regions 15 corresponding to mainly the triterpenoid region and the oligo-saccharide portions of the molecules, respectively. The exact amounts of sugars can not be determined at this point.

c)  $^1\text{H}$  NMR

20 Fig. 11 demonstrates the full proton spectrum (0-10 ppm) and Fig. 12 partial proton spectrum (anomeric region, 4.0-  
-6.0 ppm), respectively, of fractions: A, B2; B, B3; and C,  
25 B4B. The spectra are obtained from samples ( $\approx$  10 mg,  
 $\approx$  600 scans) dissolved in DMSO. $d_6$ /D<sub>2</sub>O (98:2, v/v). To the  
far left in the spectrum (Fig. 11), at 9.4 ppm, the signal  
from the aldehyde proton on carbon-24 (see Fig. 5) is found.  
The doublet nature of this peak, a peak which is supposed to  
be a singlet, since it has no neighbouring protons to couple  
30 to, offers an explanation to the unusually complex anomeric  
region which is poorly resolved (as seen in the expansion in  
Fig. 12).

35 The doublet can be due to an aldehyde proton in two different compounds or to the presence of chemical exchange between two different populations of the same molecule (this process is

slow enough in NMR time scale to be observed) thus explaining the different integrals of the peaks in the doublet at different temperatures as shown in Fig. 13 and Table 5.

5

Table 5

<u>Temperature in Degree K</u>	<u>Shift 1</u>	<u>Shift 2</u>	<u>Difference in Shifts</u>	<u>Integral Quote Shift 1/Shift 2</u>
10	301	9.46	9.44	0.02
	351	9.47	9.46	0.01
	361	9.48	9.47	0.01

Fig. 13 and Table 5 (above) demonstrate that the relative integral of the peaks varies with temperature and that the two peaks move closer to each other at a higher temperature, both indicating that it can not be two different molecules but rather two different populations of the same molecule. This would explain the complex anomeric region by suggesting that many anomeric protons in the molecule would have double resonances due to different chemical environments in the two populations. However, the present set of data indicates that differences in the glycosylation of the compounds could provide part of the explanation of their structural differences (Fig. 12), by demonstrating different amount of anomeric proton signals in the spectra. The FAB-MS data for fractions B2, B3 and B4B also does not rule out the formal possibility that two similar size molecules, with very similar physico-chemical properties, exist that have the same amount of sugars but differ in linkage-positions and/or sequence.

30

Conclusion: In general, 1-dimensional  $^1\text{H}$  NMR spectra from 8-10 sugar containing earlier unknown molecules are not sufficient for assignment of protons and detailed structural characterization. For resolving all signals and for making proper assignments through out the compounds it will be necessary to use the 2-dimensional NMR technique as well as

chemically degrade the compounds for analysis. Both homo-nuclear ( $^1\text{H}$ - $^1\text{H}$ ) and heteronuclear ( $^1\text{H}$ - $^{13}\text{C}$ ) COSY, TOCSY as well as NOESY. The 2-dimensional proton phase sensitive correlation double quantum filtered NMR spectrum (DQFPSCOSY) 5 for fraction B3 is shown in Fig. 14.

d) Summary of Structural Data

The conclusion of data generated thus far is that the active 10 fractions that have adjuvant activity and ISCOM particle forming capacity in Quil A contain unique glycosylated triterpenoid-saponins that differ between each other in both their triterpenoid and glycan parts. They have an approximate structure like the one described in Fig. 10 and consist of a 15 five-ring steroid skeleton of  $\beta$ -amyrin type and contains 8-11 sugar residues.

Example 7

0.1 mg cholesterol was mixed with  $^3\text{H}$ -cholesterol (10 mg/ml 20 dissolved in 20% MEGA-10 in  $\text{H}_2\text{O}$ ) and 0.5 mg B2 or B3 or B4B or mixtures thereof. The volume was adjusted to 0.5 ml and the mixture dialysed against PBS on a preparation treated with ammonium molybdate (negative colouring technique). The dialysed preparations were analysed for the presence of complex with 25 iscom structure by electron microscopy (EM) and analytical gradient centrifugation. In EM the iscom structure is characterized by a cage-like particle with a diameter of 40 nm composed of subunits with annular structure with a diameter of 12 nm. For sedimentation studies the sample is placed over a 30 sucrose gradient (10-50%) and centrifuged for 18 hours, +10°C in a TST 41,14 rotor, 40 000 rpm. The gradient is collected in 18 fractions (fraction 1 = the bottom and fraction 18 = the top). By localizing the  $^3\text{H}$ -cholesterol activity in the gradient, one can tell the sedimentation constant 35 and see if complexes have been made.

B4B forms typical iscom structures with cholesterol but has no potent adjuvant activity.

5 B2 does not form iscom-like structures with cholesterol but binds to cholesterol. Together with B4B, B2 forms iscom-like structures with cholesterol. B2 has a weak adjuvant activity.

10 B3 binds to cholesterol but not in iscom-like structures. With B4B, B3 like B2, can form iscom-like structures with cholesterol. B3 has adjuvant activity.

CLAIMS

1. A matrix which is not a lipid vesicle and without any intentional antigens or antigenic determinants comprising at least one lipid and at least one saponin for use as an immunomodulating agent.

5

2. A matrix according to Claim 1, comprising at least one lipid, at least one saponin and at least one adjuvant.

10 3. A matrix according to Claim 2, comprising a sterol, preferably cholesterol, one or more saponins, one or more adjuvants and one or more further lipids.

15 4. A matrix according to any of Claims 1-3, characterized in that the saponin is a triterpensaponin, especially Quil A or one or more components thereof.

5. A matrix according to Claims 1-4, comprising one or more immunomodulating compounds.

20 6. A matrix according to any of Claims 1-5, characterized in that it has an open spherical structure consisting of circular sub-units or parts of the spherical structure under electron microscopy and a sedimentation constant of about 12-22 S.

25 7. A process for preparing a matrix which is not a lipid vesicle and without any intentional antigens or antigenic determinants comprising at least one lipid and at least one saponin and optionally an adjuvant for use as an immunomodulating agent or a vaccine, characterized in that at least one lipid is solubilized in a solvent, the saponin or saponins are added, the other adjuvants and lipids are optionally also added, whereafter the solvent may be removed for example by dialysis, ultra filtration, gel filtration or electrophoreses, or be diluted.

8. A process according to Claim 7, characterized in that the lipid is a sterol, especially cholesterol, and the saponin is a triterpensaponin, especially Quil A or one or more components thereof.

5

9. A process according to Claim 7 or 8, characterized in that one or more immunomodulating compounds are added.

10

10. A process according to Claim 7-9, characterized in that the sterol and saponin are solubilized in the lipids and/or adjuvants.

15

11. Glycosylated triterpenoid saponins derived from Quillaja Saponaria Molina of Beta Amyrin type with 8-11 carbohydrate moieties which have the following characteristics:

a) Substance B2 has a molecular weight of 1988, a carbon 13 nuclear magnetic resonance (NMR) spectrum as indicated on Fig.:s 5A and 6A and a proton NMR spectrum as shown in Fig.:s 11A and 12A;

20

b) Substance B3 has a molecular weight of 2150 and has a carbon 13 NMR spectrum as shown in Fig.:s 5B and 6B, and a proton NMR spectrum as shown in Fig.:s 11B and 12B;

c) Substance B4B has a molecular weight of 1862, a carbon 13 NMR spectrum as shown in Fig.:s 5C and 6C, and a proton NMR structure as shown in Fig.:s 11C and 12C.

25

12. A process for preparing glycosylated triterpenoid saponins derived from Quillaja Saponaria Molina of Beta Amyrin type with 8-11 carbohydrate moieties which have the following characteristics:

30

a) Substance B2 has a molecular weight of 1988, a carbon 13 nuclear magnetic resonance (NMR) spectrum as indicated on Fig.:s 5A and 6A and a proton NMR spectrum as shown in Fig.:s 11A and 12A;

35

b) Substance B3 has a molecular weight of 2150 and has a carbon 13 NMR spectrum as shown in Fig.:s 5B and 6B, and a proton NMR spectrum as shown in Fig.:s 11B and 12B;

c) Substance B4B has a molecular weight of 1862, a carbon 13 NMR spectrum as shown in Fig.:s 5C and 6C, and a proton NMR structure as shown in Fig.:s 11C and 12C, characterized in extracting the bark from Quillaja Saponaria Molina with water, dialysing the aqueous extract, lyophilizing the dialysate, subjecting the lyophilized matter to gel filtration on Sephadex G-50 and subsequent weak anionic exchange chromatography on DEAE-cellulose, eluting the products with 0.2M sodium chloride at neutral (pH) conditions and separating them by repeated chromatography on silica gel.

13. A vaccine comprising a matrix as hereinbefore defined according to Claims 1-6, one or more antigens and a pharmaceutically acceptable vehicle.

14. A kit for human or veterinary medicine use, characterized in that it comprises at least one matrix according to Claims 1-6 and one or more immunomodulating substances and a pharmaceutically acceptable vehicle in mixture or in separate compartments.

1/11

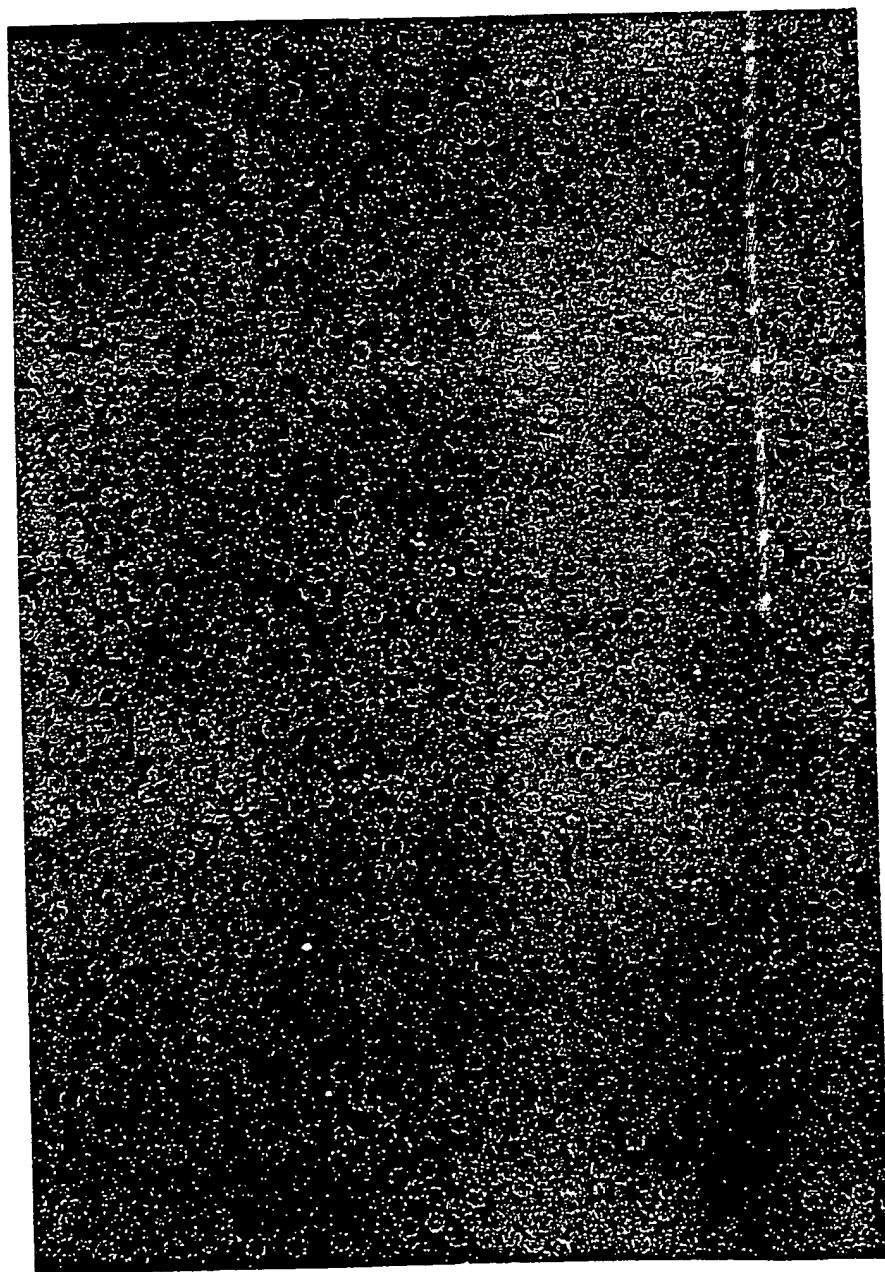


FIG. 1

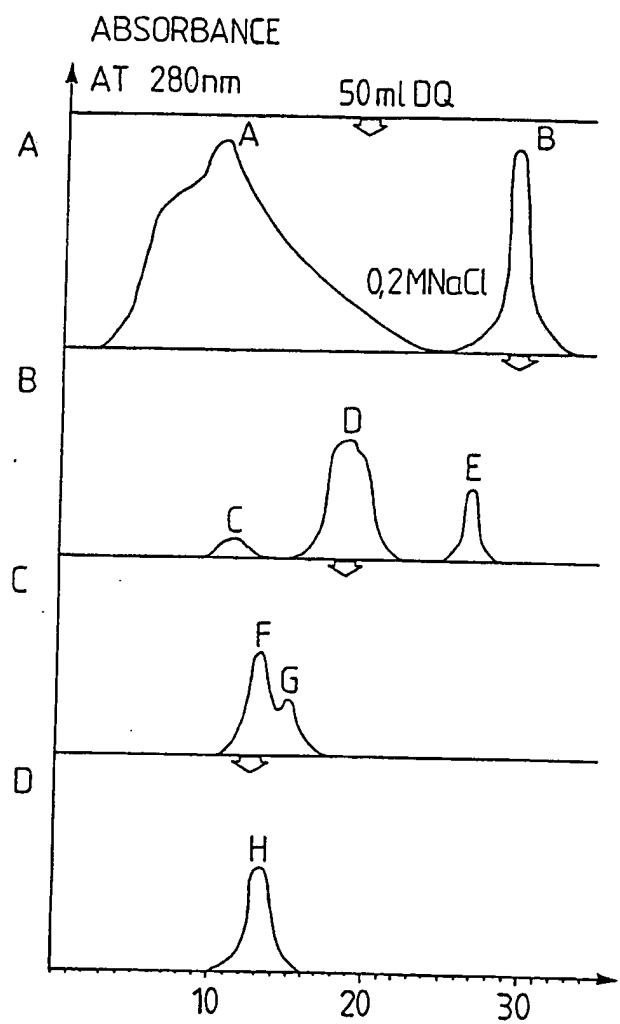


FIG.2

3/11

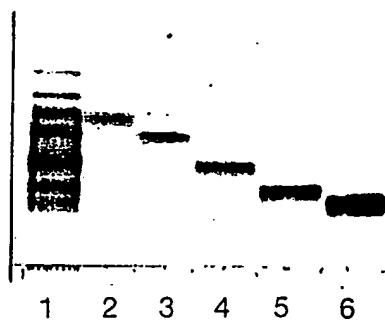


FIG. 3

4/11

## FAB-SPECTRA

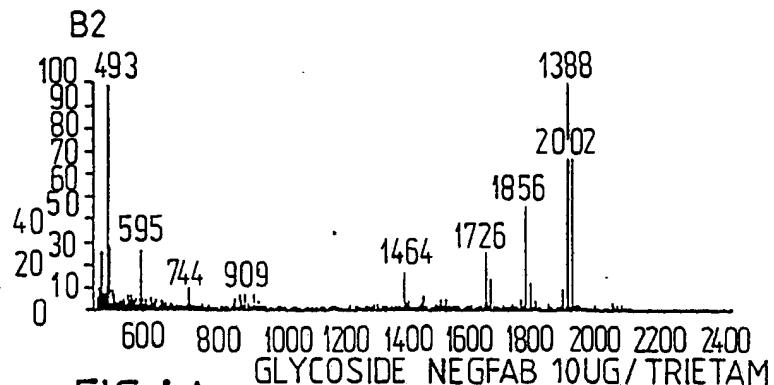


FIG.4A

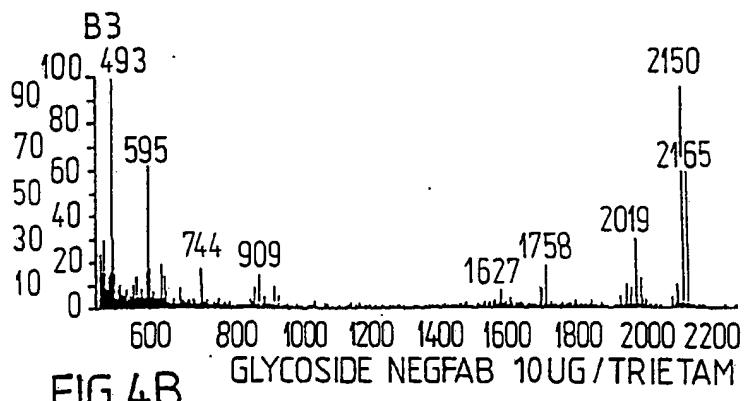


FIG.4B

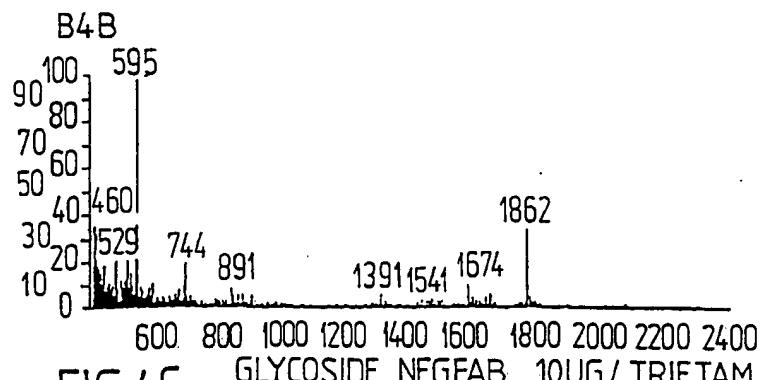


FIG.4C

5/11



FIG.5A

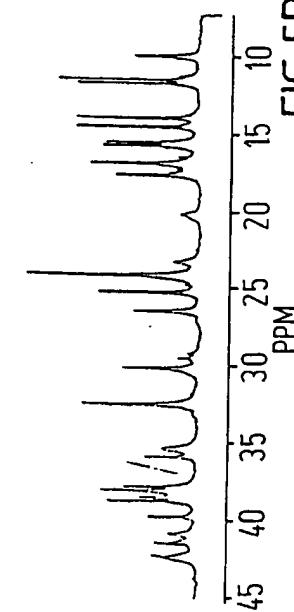


FIG.5B

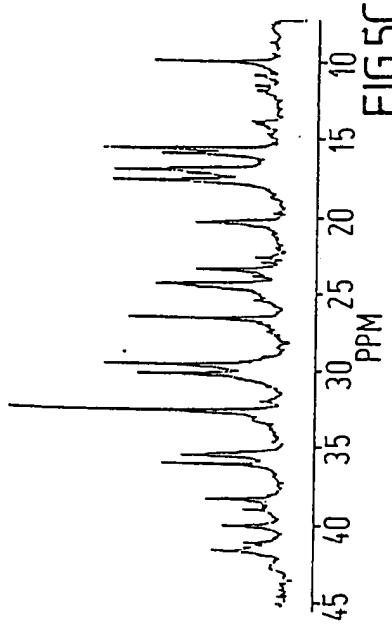


FIG.5C

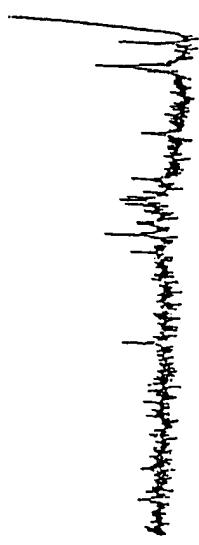


FIG.6A

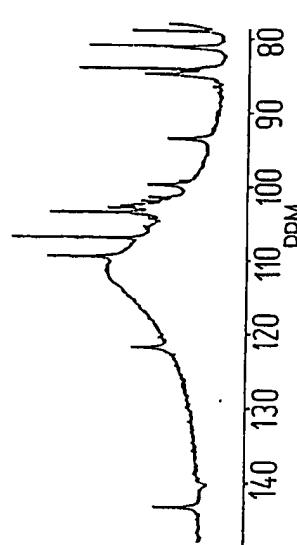


FIG.6B

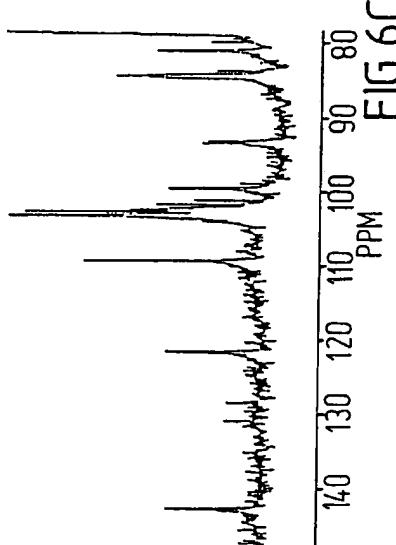
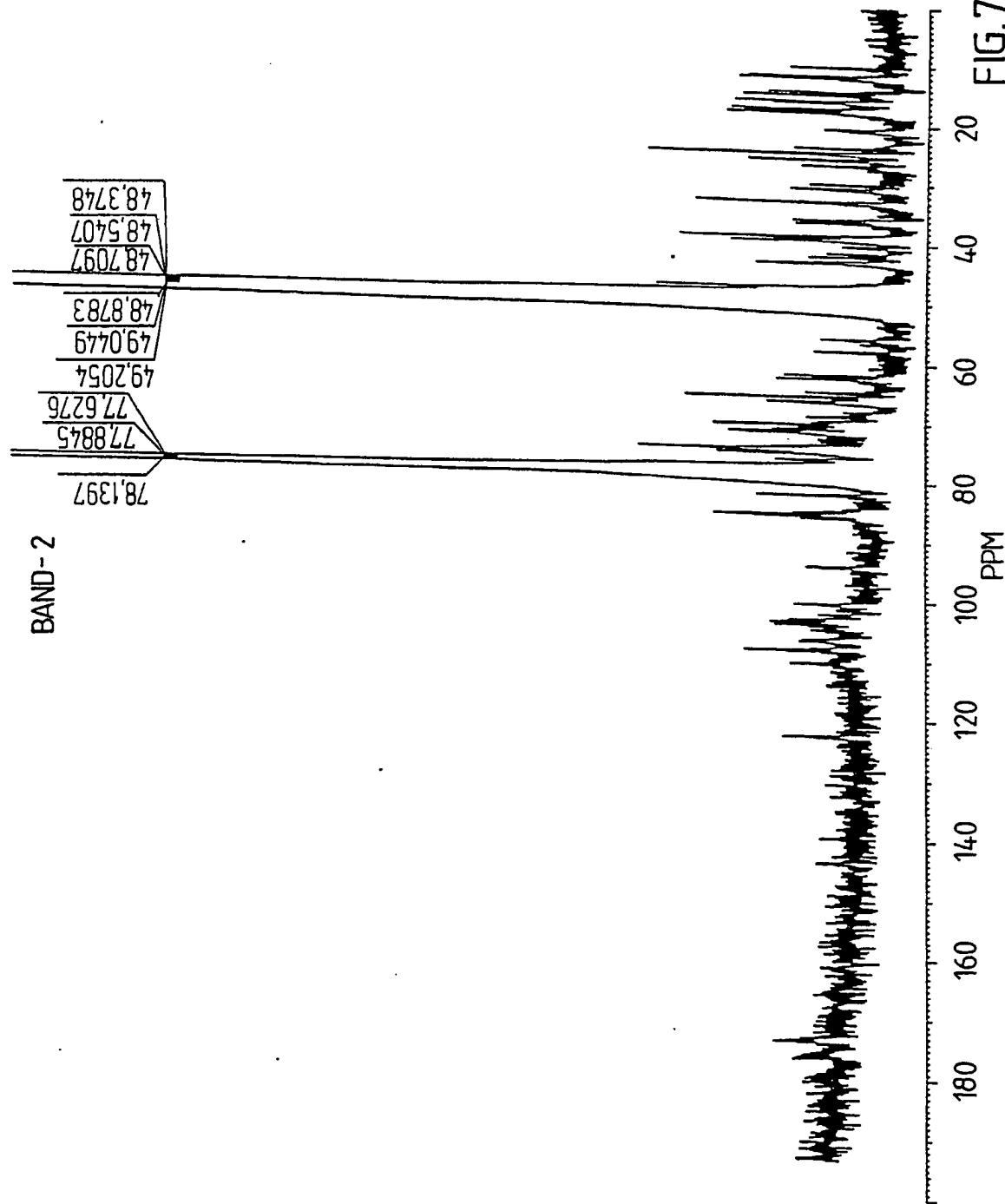
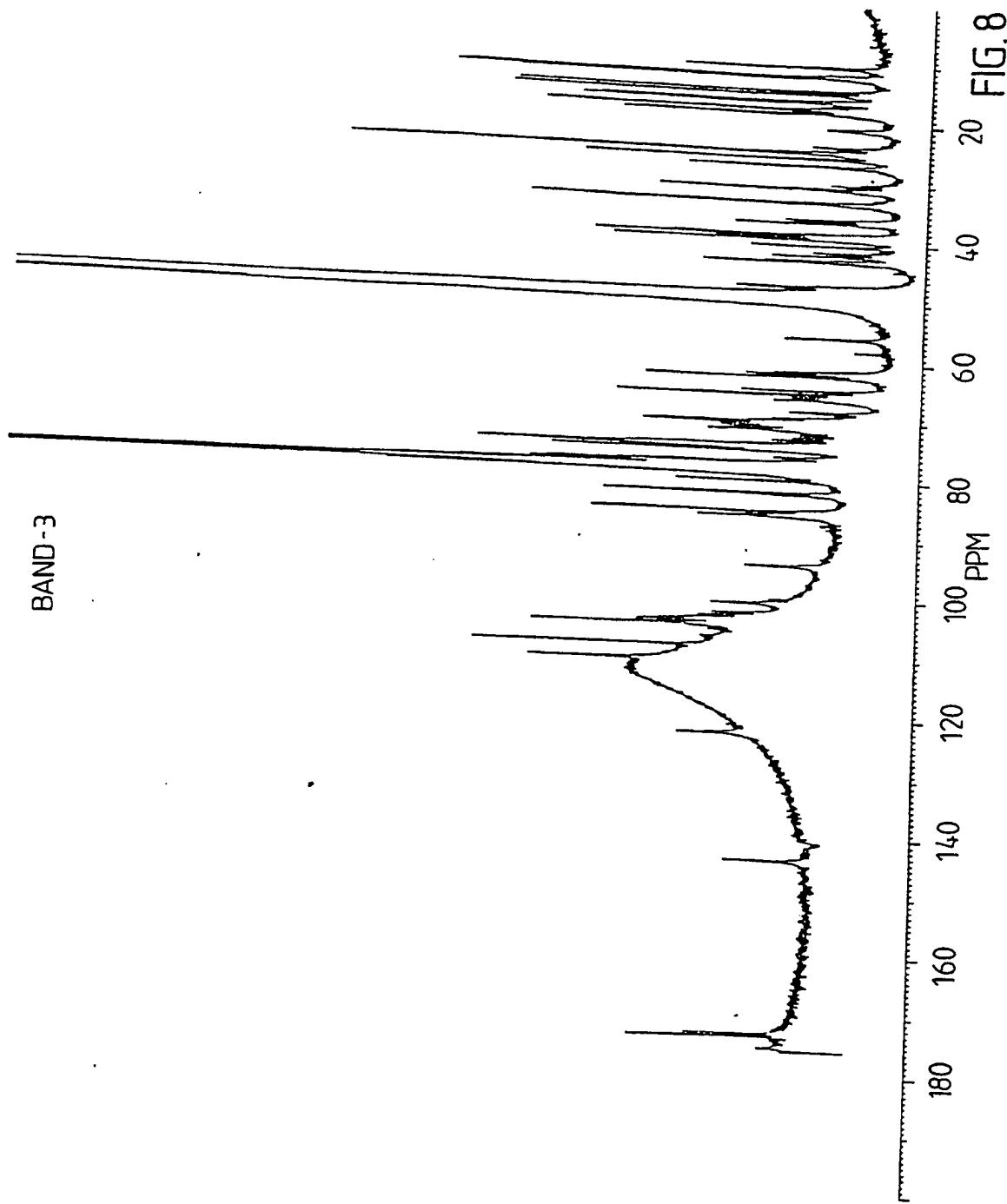


FIG.6C

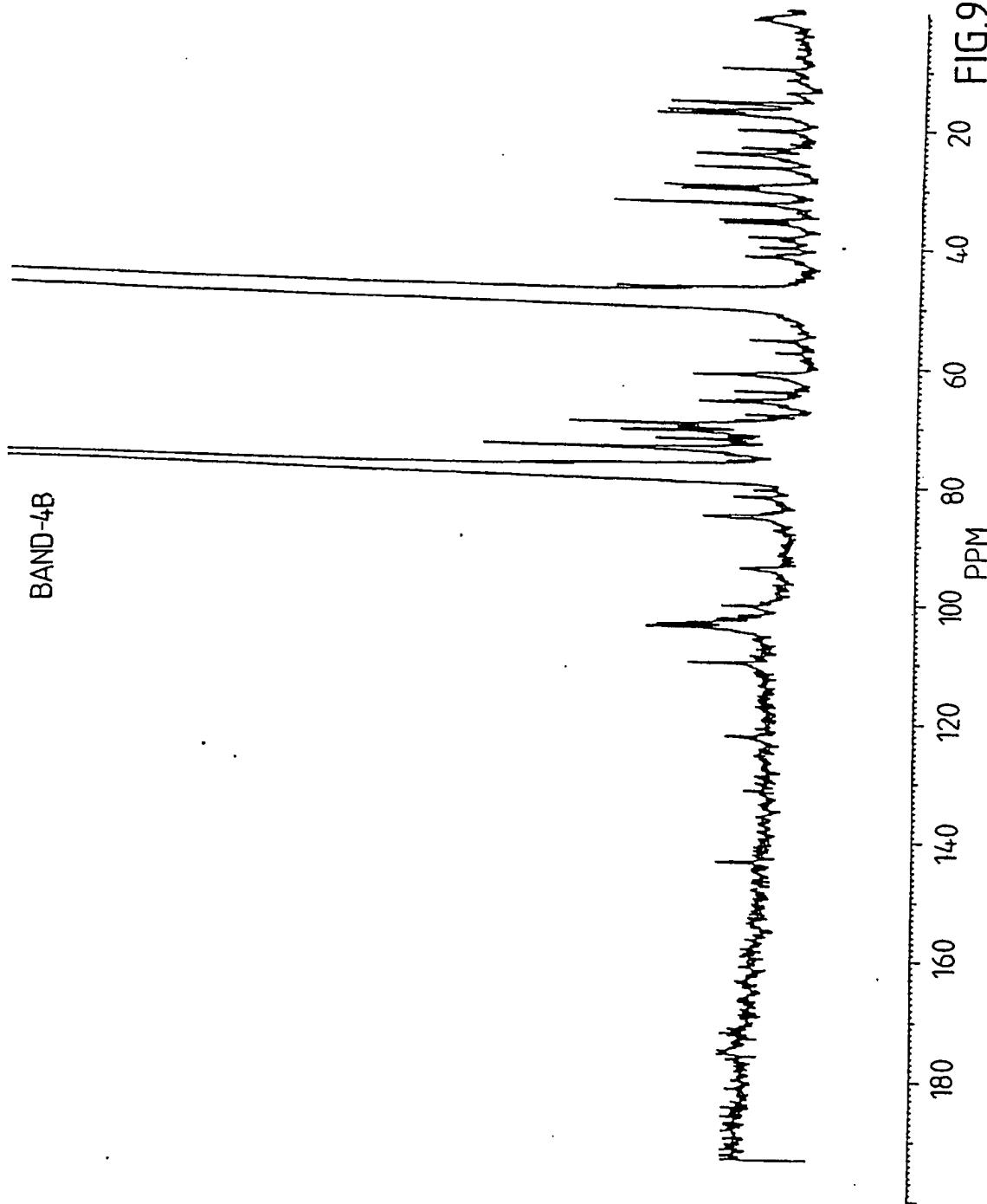
6/11

**SUBSTITUTE SHEET**

7/11



8/11



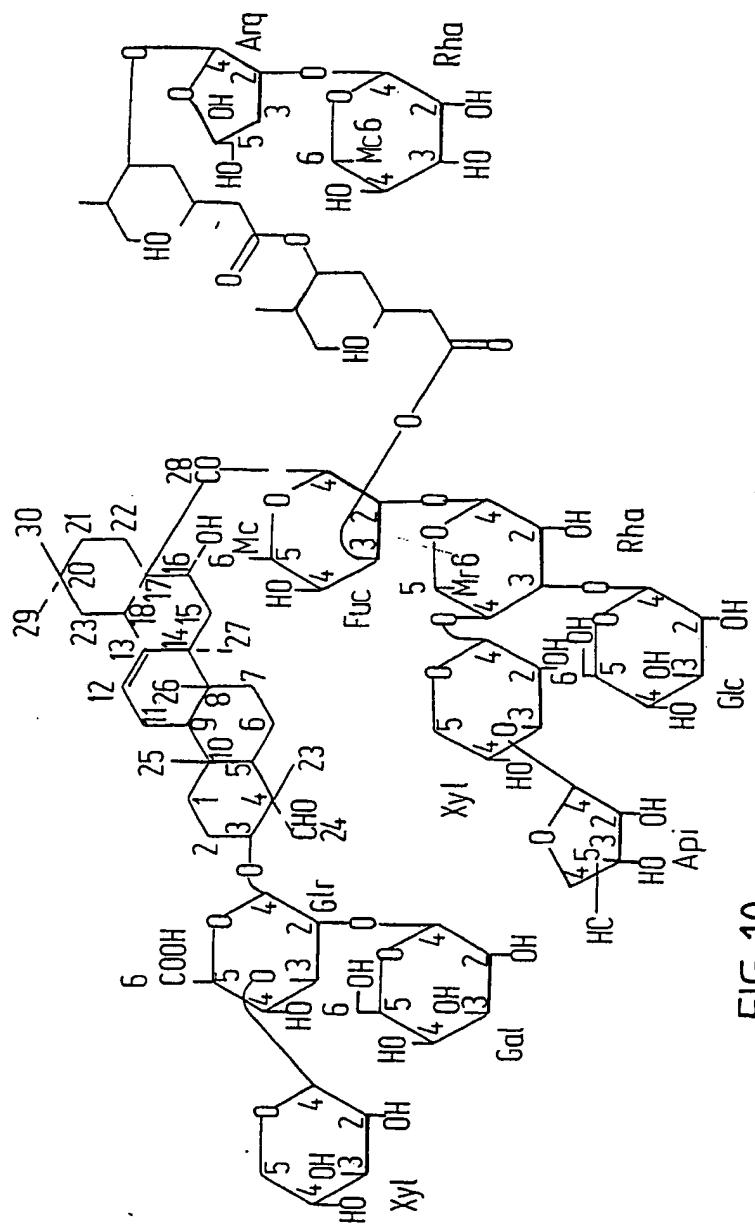


FIG. 10

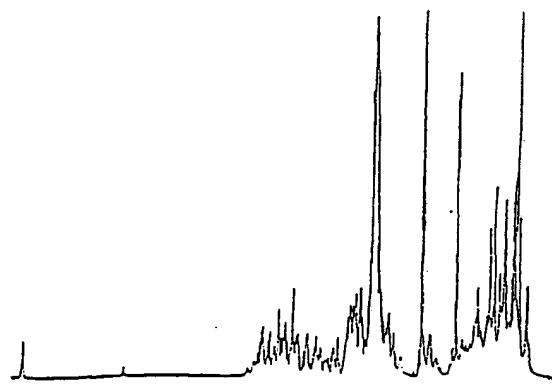


FIG. 11A PPM

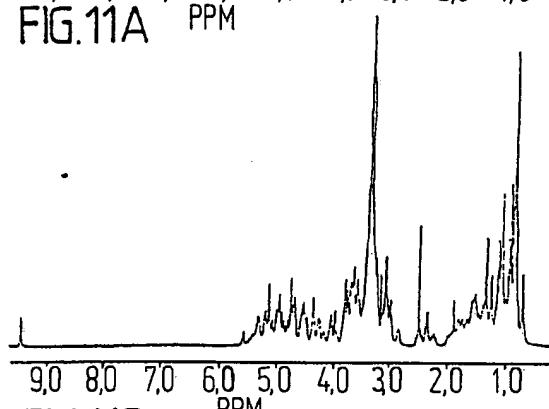


FIG. 11B PPM

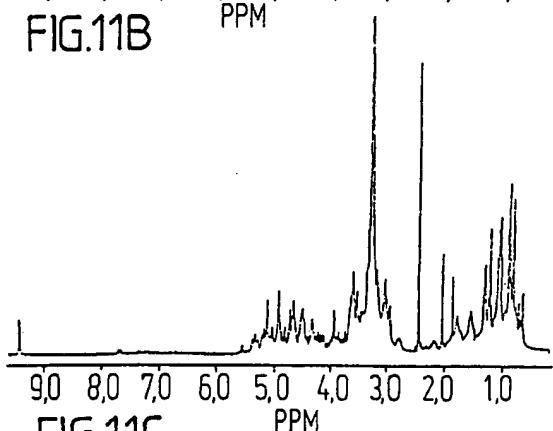


FIG. 11C PPM

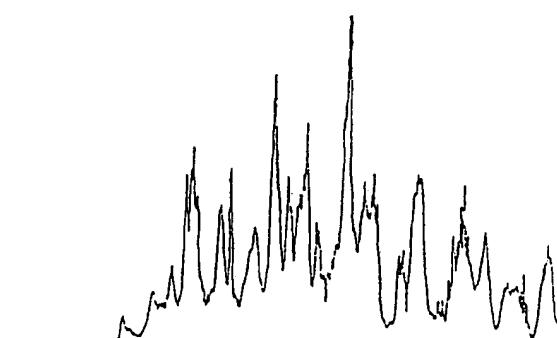


FIG. 12A PPM

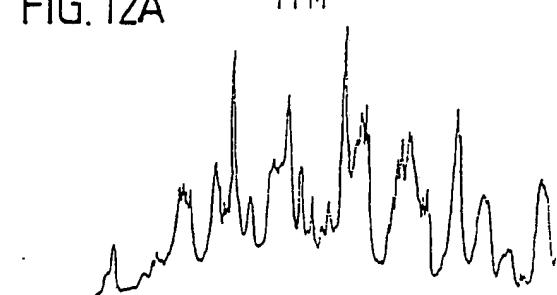


FIG. 12B PPM

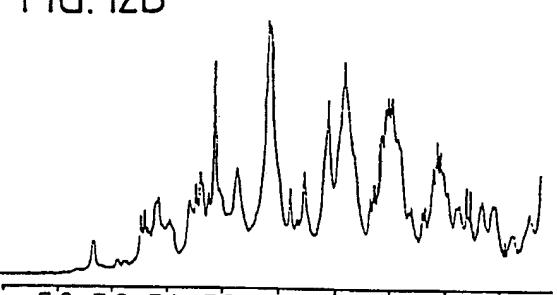


FIG. 12C PPM

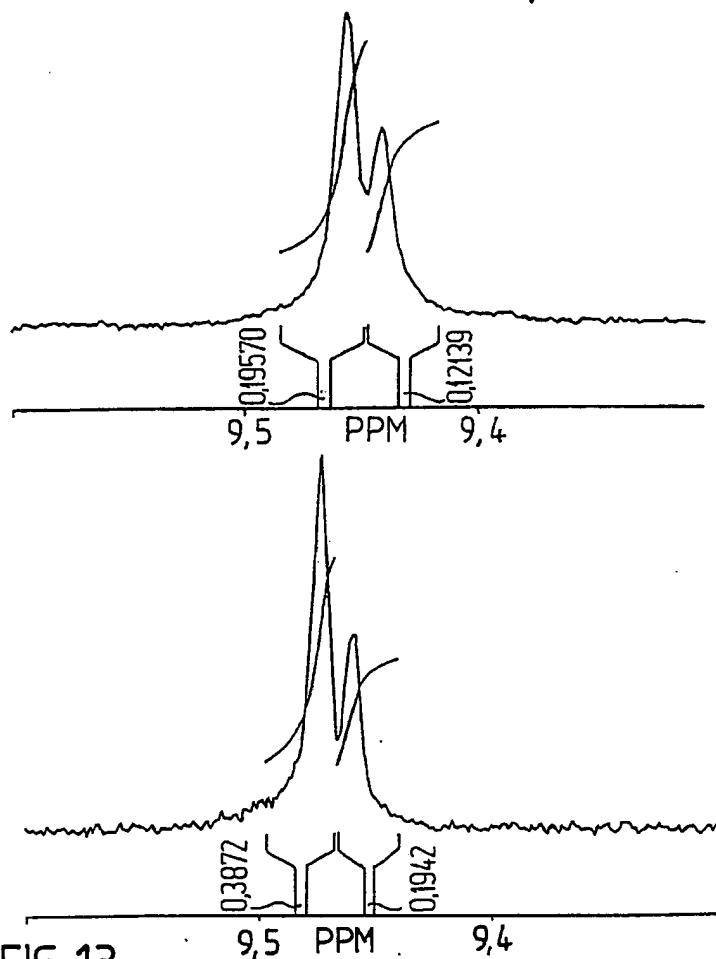


FIG. 13

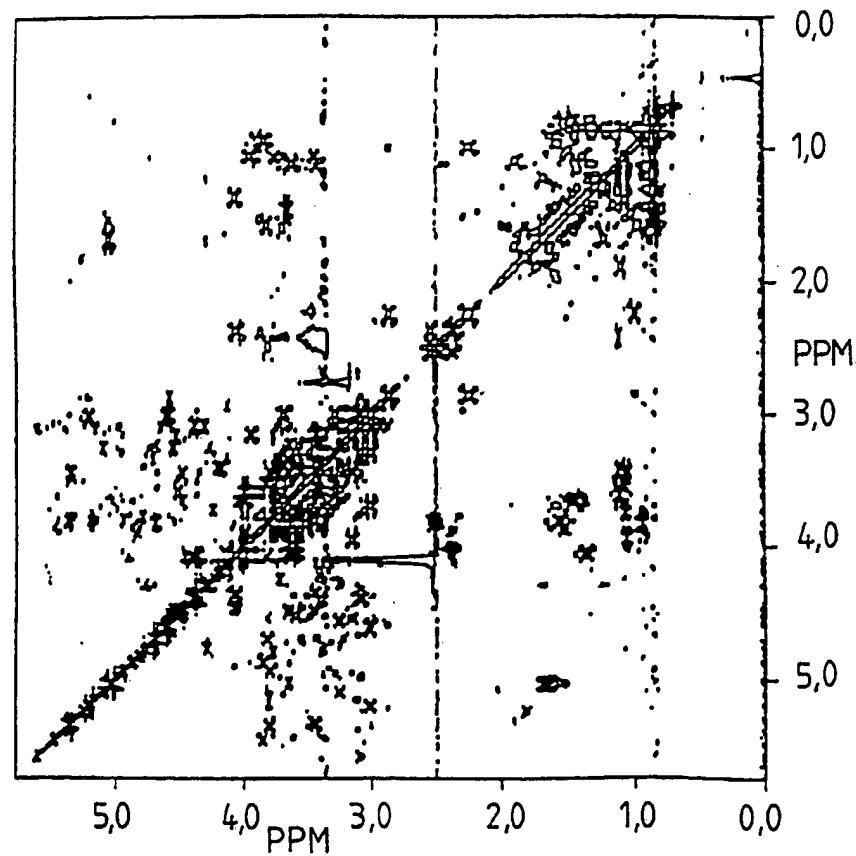


FIG. 14

SUBSTITUTE SHEET

# INTERNATIONAL SEARCH REPORT

International Application No. PCT/SE 89/00528

## I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) <sup>6</sup>

According to International Patent Classification (IPC) or to both National Classification and IPC  
 IPC4: A 61 K 39/39, C 07 J 17/00, 63/00

## II. FIELDS SEARCHED

Minimum Documentation Searched <sup>7</sup>

Classification System	Classification Symbols
IPC4	CA Search A 61 K; C 07 J

Documentation Searched other than Minimum Documentation  
to the Extent that such Documents are Included in the Fields Searched <sup>8</sup>

SE,DK,FI,NO classes as above

## III. DOCUMENTS CONSIDERED TO BE RELEVANT <sup>9</sup>

Category <sup>10</sup>	Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>	Relevant to Claim No. <sup>13</sup>
X	Int. Archs Allergy appl. Immun., Vol. 67, 1982 R. Bomford: "Studies on the Cellular Site of Action of the Adjuvant Activity of Saponin for Sheep Erythrocytes ", see page 127 - page 131 see esp. p. 130	1
Y	--	1
Y	Biotechnology and applied Biochemistry, Vol. 10, 1988 K. Lövgren et al.: "The Requirement of Lipids for the Formation of Immunostimulating Complexes (Iscoms) ", see page 161 - page 172 see esp. p. 164-167	1
X	EP, A1, 0 231 039 (DE STAAT DER NEDERLANDEN VERTEGENWOORDIGD DOOR DE MINISTER VAN WELZIJN, VOLKSGEZONDHEID EN CULTUUR) 5 August 1987, see the whole document	7
Y	--	1

\* Special categories of cited documents: <sup>10</sup>

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

## IV. CERTIFICATION

Date of the Actual Completion of the International Search 12th December 1989	Date of Mailing of this International Search Report 1989-02-01
International Searching Authority SWEDISH PATENT OFFICE	Signature of Authorized Officer Carl Olof Gustafsson

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No
P,X	WO, A1, 88/09336 (CAMBRIDGE BIOSCIENCE CORPORATION) 1 December 1988, see esp. p. 6-8 --	11
A	US, A, 4 806 350 (JAY D. GERBER) 21 February 1989, see the whole document --	1-5
A	US, A, 4 101 652 (A. BONATI) 18 July 1978, see the whole document --	1
A	Int. Archs Allergy appl. Immun., Vol. 63, 1980 R. Bomford: "Saponin and Other Haemolysins (Vitamin A, Aliphatic Amines, Polyene Antibiotics) as Adjuvants for SRBC in the Mouse", see page 170 - page 177 --	1-5
A	Infection and Immunity, Vol. 56, No. 2, February 1988 Gideon F.A. Kersten et al.: "Incorporation of the Major Outer Membrane Protein of Neisseria gonorrhoeae in Saponin-Lipid Complexes (Iscoms): Chemical Analysis, Some Structural Features, and Comparison of Their Immunogenicity with Three Other Antigen Delivery Systems", see page 432 - page 438 --	1-5
A	Archiv für die gesamte Virusforschung, Vol. 44, 1974 K. Dalsgaard: "Saponin Adjuvants ", see page 243 - page 254 --	12
X	Chemical Abstracts, volume 108, no. 20, 16 May 1988, (Columbus, Ohio, US), see page 398, abstract 173547k, & JP,, 62205025 (Triterpenoid saponins as antiulcer agents.) 9 September 1987 -----	11

## FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V.  OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE<sup>1</sup>

This International search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1.  Claim numbers ..... because they relate to subject matter not required to be searched by this Authority, namely:

2.  Claim numbers ..... because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3.  Claim numbers....., because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI.  OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING<sup>2</sup>

This International Searching Authority found multiple inventions in this international application as follows: The claims 1-14 define two independent inventions, namely: Claims 1-10, 13-14 drawn to a matrix, comprising at least one lipid and at least one saponin, a process for preparing the matrix, a vaccine comprising the matrix and a kit for medicine use. Claims 11-12 drawn to specified glycosylated triterpenoid saponins and a process for preparing those.

1.  As all required additional search fees were timely paid by the applicant, this International search report covers all searchable claims of the international application.

2.  As only some of the required additional search fees were timely paid by the applicant, this International search report covers only those claims of the international application for which fees were paid, specifically claims:

3.  No required additional search fees were timely paid by the applicant. Consequently, this International search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

4.  As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

## Remark on Protest

- The additional search fees were accompanied by applicant's protest.  
 No protest accompanied the payment of additional search fees.

**ANNEX TO THE INTERNATIONAL SEARCH REPORT  
ON INTERNATIONAL PATENT APPLICATION NO. PCT/SE 89/00528**

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.  
**08/11/89**

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP-A1- 0 231 039	05/08/87	JP-A-	63002933	07/01/88
WO-A1- 88/09336	01/12/88	AU-D-	19340/88	21/12/88
US-A- 4 806 350	21/02/89	EP-A-	0242205	21/10/87
		AU-D-	71760/87	22/10/87
		JP-A-	62255436	07/11/87
US-A- 4 101 652	18/07/78	BE-A-	843552	18/10/76
		NL-A-	7607272	04/01/77
		FR-A-B-	2315939	28/01/77
		LU-A-	75263	18/02/77
		DE-A-C-	2628143	27/01/77
		GB-A-	1503388	08/03/78
		JP-A-	52018809	12/02/77
		CA-A-	1068604	24/12/79
		CH-A-	621065	15/01/81